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(56) Documents Cited

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WO 92/22635 A1

Transgenic Research 1994, 3(3), pp.141-151 - Barash
et al.

(58) Field of Search

Online: EPODOC, WPI, BIOSIS, MEDLINE,
CAS-ONLINE, SCISEARCH, EMBASE

(54) Abstract Title

Human serum albumin expression and use in gene therapy

(57) The present invention relates to DNA vectors as a means of gene therapy for the use in treatment of human and animal disease states that are characterised by failure or inadequate functioning of the liver. This results in the lowering of the blood levels of certain important proteins produced by the liver leading to hypoalbuminaemia, anaemia, thrombocytopenia and coagulation disorders related to liver failure. The invention also relates to an expression cassette for expressing a human serum albumin into a mammalian cell comprising:

(i) a cDNA sequence of the human serum albumin,

(ii) a transcriptional control sequence operably linked to said cDNA sequence.

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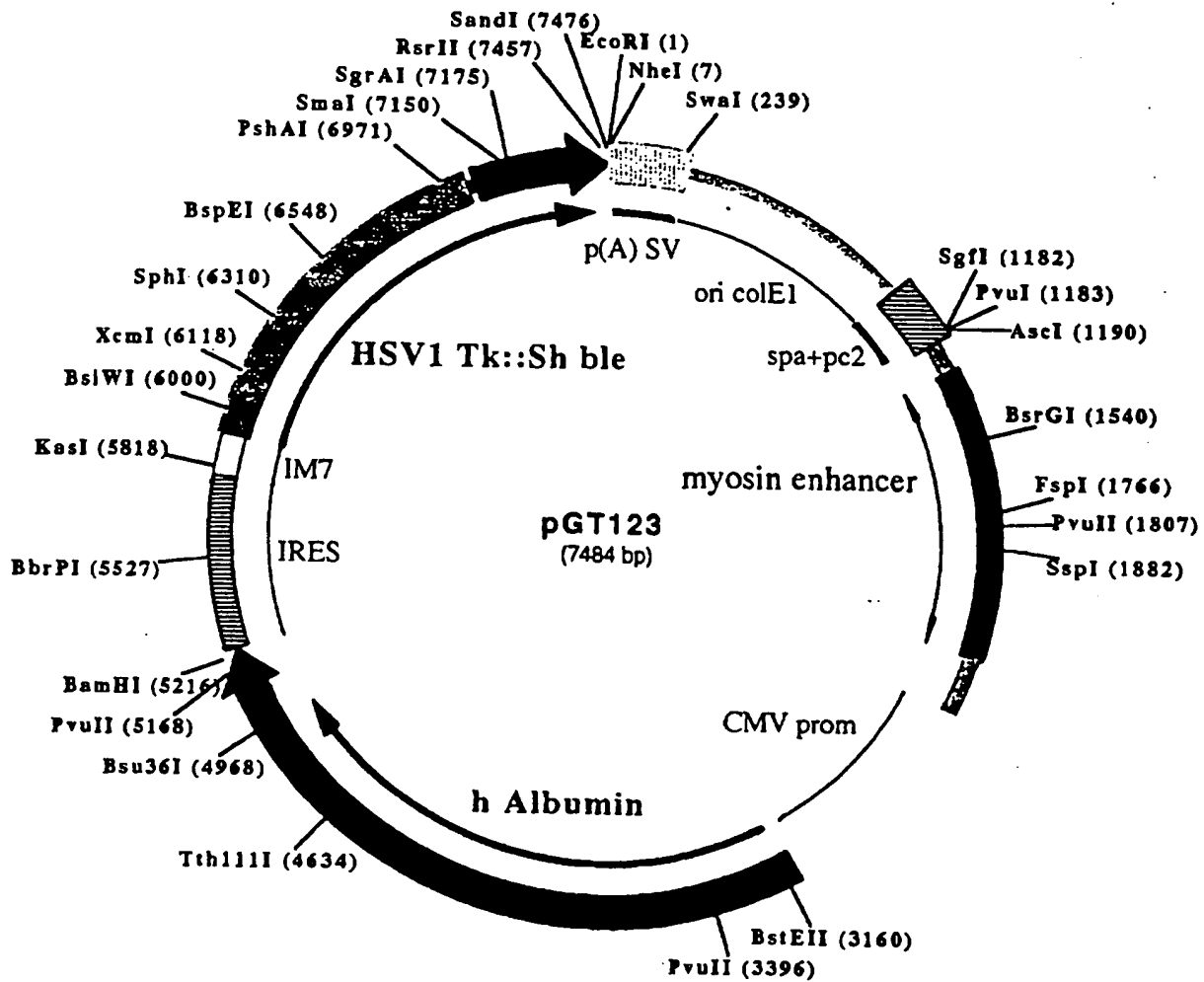
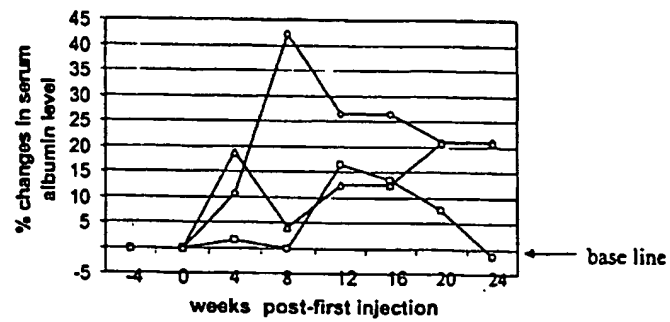


Fig. 1

Patients treated with CMV-albumin plasmid



Patients treated with myosin-albumin plasmid

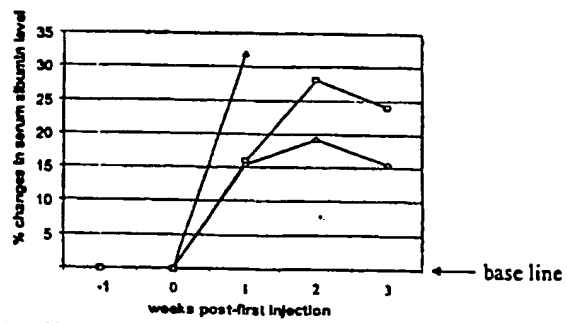


Fig 2

Changes in serum albumin level

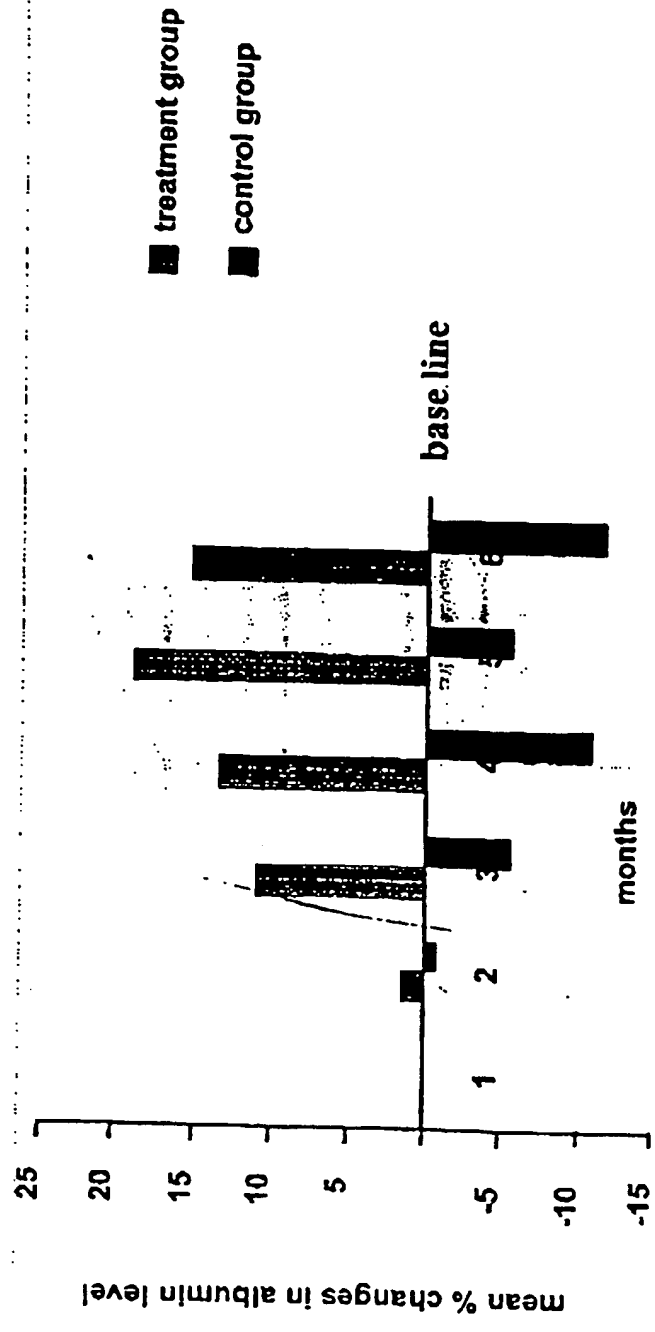


Fig 3

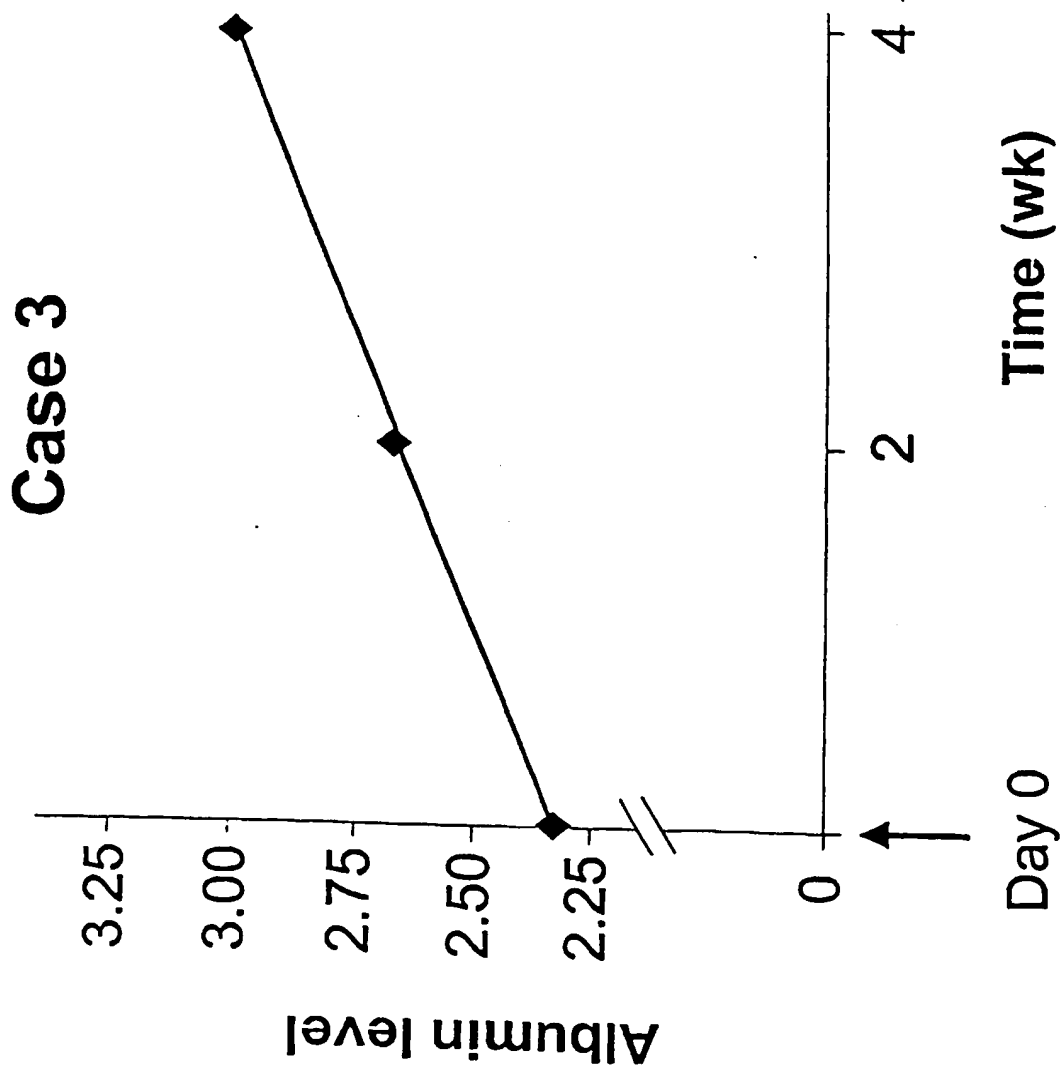


Fig 4

A METHOD FOR PROMOTING EXTRA-HEPATIC PRODUCTION OF PROTEINS FOR THE CORRECTION OF HYPOALBUMINAEMIA, ANAEMIA, THROMBOCYTOPENIA AND/OR COAGULATION DISORDERS RELATED TO LIVER FAILURE.

The present invention relates generally to the fields of human treatment of complications and side-effects disorders associated with a failure of the liver to produce proteins and albumin in particular, and more specially to expression cassettes and related recombinant DNA compounds useful in correction of low serum albumin (hypoalbuminaemia) by gene therapy.

Human serum albumin (HSA) is a naturally-occurring, globular, nonglycosylated polypeptide (MW 65,000) comprising nearly 600 amino acid residues (see e.g. Lawn et al., 1981, *Nucleic Acid Research*, 9, 6103-6113) synthesized in the liver of higher species. This protein plays important roles in the organism as it the maintenance of osmotic pressure; is involved in the binding and transport of a wide variety of substances such as fatty acids, ions (copper, nickel, calcium), bile, bilirubin, protoporphyrin, prostaglandins, steroid hormones, thyroxine, cystine, glutathione, various drugs and water soluble vitamins and the like; and additionally provides a direct source of amino acids to tissues.

In healthy conditions HSA is the most abundant protein found in the plasma of adult humans where its concentration is about 40mg/ml. However, this level can dramatically decline in the context of many diseases, especially those affecting the liver, such as for example hepatitis, cirrhosis, primary and secondary liver cancers, or infectious and inflammatory conditions, chemotherapy and radiotherapy treatments, malabsorption and nephrotic syndromes (for a medical revue, see Doweiko et al., 1991, *J. Parenteral and Enteral Nutrition*, 15, 476-483) which are generally associated with a reduced synthesis of albumin or an excessive metabolism of said protein leading to plasma low concentrations and to complications or side effects such as hypovolemia, hypoproteinemia, ascites formation, oedema, etc... Moreover, Goldwasser and Feldman (1997, *J. Clin. Epidemiol*, 50, 693-703) have reported an association between hypoalbuminaemia and mortality risk. These authors have particularly shown an increase in the odds of death ranges from 24% to 56% for each 2.5mg/ml decrement in serum albumin concentration. HSA concentrations below 35 mg/ml indicates a pathological status. The method described in the invention is a platform technology with applications for the treatment and correction of various human and animal disease states. Hypoalbuminaemia is not a single disease, but can occur as a consequence of many diseases. Therefore the invention is not restricted in its use to the treatment of a single condition. Hypoalbuminaemia can occur if the liver has an underlying disease state and is failing to produce adequate amounts of albumin. This occurs in diseases such as hepatitis, cirrhosis and liver cancer (both primary and secondary cancers). Hypoalbuminaemia also occurs as the result of

any disease state that affects the body's metabolism, such as advanced cancer, the treatment of cancer with chemotherapy or radiotherapy, catabolic states such as severe infections or inflammation, malabsorption or the nephrotic syndrome and recovery from major surgery. All of these conditions, and many others, can result in a reduced level of albumin in the blood and would be suitable for treatment using the method described in this invention. In addition, the invention could be used to provide a source of other proteins, in addition to albumin, produced at sites other than the physiological source of the proteins in the body. In this invention the prime extra site of production is the skeletal muscle, but other sites could be included. The other proteins that could be used in this invention include, but are not restricted to, erythropoietin for the correction of anaemia, thrombopoietin for the correction of thrombocytopenia, protein C and protein F for patients requiring anticoagulation, and leptin for the treatment of obesity. In addition, the invention can be used to produce hormones and growth factors used by the human body at sites other than the physiological source of such hormones and growth factors. These would include but not be restricted to insulin for the treatment of diabetes, endothelial nitric oxide synthetase (ENOS) for the prevention of thrombosis after coronary artery surgery or coronary balloon dilatation or myocardial ischaemia, Granulocyte Colony Stimulating Factor (GCSF), Macrophage Colony Stimulating Factor (MCSF) or Granulocyte Macrophage Colony Stimulating Factor (GMCSF), and/or interferon for the treatment of hepatitis or cancer.

Careful treatment of hypoalbuminaemia is a key to improving quality of life and to reducing mortality risk of affected patients. The standard treatment of hypoalbuminaemia is the administration of albumin employed as a therapeutic drug.

First, commercial HSA has been prepared from human plasma as a by product from the fractionation of donated blood. However, availability of human plasma is limited and careful heat treatment of the product prepared from human plasma should be effected to avoid potential contamination of the HSA product by hepatitis ~ and HIV viruses. Moreover, said isolation of HSA from natural sources is technically difficult, expensive and time consuming.

Against this background related to consideration of ethics, safety and stability of supply, further efforts have centered on the development of efficient recombinant methods for the production of recombinant HSA in bacterial cells or yeast which could be used as a substitute for natural HSA (Hone et al., 1998, Gen. Pharmac., 31, 811-815).

Although effectiveness of the exogenous albumin supply to the patient has been supported, indicating that serum level can be transiently restored, said treatment still means that repeated administration of albumin is required every 48 hours throughout the lifetime of the patient.

Accordingly, the prior art is deficient in providing a satisfactory treatment method providing an acceptable level of serum albumin and being compatible with quality of life of the patients. The present invention fulfills this longstanding need and desire in the art.

Since the discovery that skeletal muscle can be transfected *in vivo* by transmuscular injection of plasmid DNA, this organ system has attracted considerable attention as a potential source of secreted therapeutic proteins. However, the efficiency of this method of transfection is still low, even with the induction of muscle degeneration and regeneration through injection of myotoxic substances prior the injection of DNA. Accordingly, most studies so far have shown that expression is not high enough to increase the blood levels of circulating proteins, especially in case where this level should be high enough for permitting an improvement of the health.

It was now surprisingly found that the transfection of a polynucleotide comprising the cDNA sequence of human serum albumin into vertebrate tissue leads to a dramatic expression of HSA which provides an efficient tool for correcting hypoalbuminaemia and associated disorders. In particular, it was surprisingly found that injection into muscular tissue of said polynucleotide leads to a significant augmentation of serum albumin levels for up to three months and to reduced complications or side effects in treated patients. Thus, the present invention first relates to an expression cassette, reliable recombinant DNA compound and their use for the preparation of a pharmaceutical composition which provide a steady constitutive level of expression allowing an effective amount of protein to be produced and a restoring in the treated patients albumin levels leading to improved clinical status. Generally, the present invention provides a gene therapy approach aiming to increase the serum albumin level at least to the lower value of the normal range.

Therefore, the technical problem underlying the present invention is the provision of improved methods and means for the correction of hypoalbuminaemia by gene therapy. This technical problem is solved by the provision of the embodiments as defined in the claims.

The present invention concerns an expression cassette for expressing a human serum albumin into a mammalian cell comprising:

- (i) a cDNA sequence of the human serum albumin,
- (ii) a transcriptional control sequence operably linked to said cDNA sequence.

According to a preferred embodiment, said expression cassette comprises a cDNA sequence of the human serum albumin having the nucleotide sequence set forth in SEQ ID NO:1

Said expression cassette can also be constructed by using a cDNA sequence of the human serum albumin of about 2.0kb in length obtained by enzymatic digestion by KpnI and SalI of the plasmid PILMALB5 having the deposit accession number HSRRB HG226.

It should be also noticed here that the cDNA production from mRNA is well described in the literature and that two cDNA sequences could be obtained from a unique mRNA depending on the oligonucleotide sequences used in order to initiate the reverse transcription of said mRNA. Said cDNA sequences differ only at their respective 5'end and/or 3'end where sequence variations can occur. Nevertheless, each of said cDNA sequences will comprise the correct sequence corresponding to said mRNA and accordingly can be used in the context of the present invention.

"operably linked" means that the cDNA and the transcriptional control sequences are in a relationship permitting them to function in their intended manner. Thus, for example, a promoter operably linked to a cDNA sequence is ligated in such a way that expression of the human serum albumin is achieved under conditions which are compatible with the transcriptional activity of the promoter. These conditions are widely used in the technical field of the invention.

"Expression of a gene, and therefore of the corresponding polypeptide", means that the encoding DNA sequence is first transcribed in mRNA, which itself is further translated in the corresponding polypeptide sequence.

"Human serum albumin or HAS" according to the present invention preferably designates a polypeptide sequence as set forth in SEQ ID N:2 (Lawn) - However, comparison analysis of published HSA sequences (Lawn et al., 1981, Nucleic Acid Res., 9, 6103-6114 and Dugaiczky et al., 1982, PNAS, 79, 71-75 for example) shows that minor amino acid variations are acceptable in the HSA polypeptide sequence which do not affect the HSA properties. Accordingly, and with regard to the degenerescence of the genetic code, the skilled man can easily adapt the cDNA sequence of the present invention with respect to these minor changes. These specific embodiments are encompassed by the invention.

"Transcriptional control sequence" designates the nucleic acid sequences which control the initiation of the transcription, the selection of the start position, which regulate the transcription level (enhancement or inhibition), which determine the type of polymerase directing the polymerisation of the transcribed mRNA, which control the transcription rate, the termination of said transcription, and the site of said termination. These sequences are widely analyzed, used and reported in the literature and can be readily obtained or adapted by those skilled in the art.

In a specific embodiment, the transcriptional control sequence according to the invention comprises a promoter element which is selected from the group consisting of viral promoters. Examples of such viral promoters are the SV40 early and late promoters, the adenovirus major late



5

promoter, the Rous Sarcoma Virus (RSV) promoter, the Cytomegalovirus (CMV) immediate-early promoter, the herpes simplex virus (HSV) promoter, MPSV promoter, 7.5k promoter, vaccinia promoter and the Major-intermediate-early (MIE) promoter. The Cytomegalovirus (CMV) immediate-early promoter is however preferred.

The promoter nucleic acid sequence can be a naturally occurring promoter isolated from biological nucleic acid material or chemically synthesised. The promoter sequence can also be artificially constructed by assembling elements previously screened for transcriptional activity leading to potencies which can exceed those of naturally occurring ones (Li et al., 1999, Nature Biotech., 17, 241-245)

According to the present invention, "nucleic acid sequence" means a fragment or a portion of a nucleic acid, without size limitation, which may be either linear or circular, natural or synthetic, modified or not (see US 5525711, US 4711955 US 5792608 or EP 302175 for modification examples). Depending on the considered sequence, it may be, *inter alia*, a genomic DNA, a cDNA or a synthesised DNA. More elaborated "nucleic acid sequences" such as "recombinant DNA compound" may be in the form of linear nucleic acid construct, and preferably in the form of plasmid. According to the invention, said "recombinant DNA compound" should preferably be understood as a naked nucleic acid construct or "plasmid vector" (Wolff et al., Science 247 (1990), 1465-1468), or as nucleic acid construct formulated with at least one compound such as polypeptides, preferably viral polypeptides (for a review, see Robbins et al., 1998, Tibtech, 16, 35-40) (said "recombinant DNA compound" is herein termed "viral vector"), or cationic lipids or cationic polymers which can participate in the uptake or stabilization of the nucleic acid construct into the cells (see Ledley, Human Gene Therapy 6 (1995), 1129-1144 or Rolland, 1998, Therapeutic Drug Carrier Systems, 15, 143-198 for a review) (said "recombinant DNA compound" is termed "synthetic vector"). According to the invention, the nucleic acid sequence can be homologous or heterologous to the host expressing cells.

The expression cassette can be constructed using routine cloning techniques known to persons skilled in the art (for example, see Sambrook et al., 1989, Molecular Cloning-a laboratory manual ; Cold Spring Harbor Press).

In still another aspect of the invention, the transcriptional control sequence further comprises at least one enhancer element. The definition of enhancers refers to regulatory elements which activate transcription in a position and orientation independent way. Several enhancer elements have been identified to date in many genes. Preferably, the present enhancer element is a myosin light chain enhancer. More preferably, the enhancer used in the expression cassette of the present invention is of vertebrate origin, more preferably of mammalian origin. The rat myosin light chain 1/3 enhancer

(Donoghue et al., 1988, Gene & Dev., 2, 1779-1790) is especially preferred. The enhancer element is operably linked to the promoter, may be localized either upstream or downstream of said promoter and may be used in either orientation. In a specific construction of the present invention, said enhancer sequence is located upstream of said cDNA sequence of the human serum albumin. According to a preferred embodiment, said transcriptional control sequence comprises at the most three enhancer sequences, the sequence of which is identical or selected independently of one another.

The present invention is more specifically directed to an expression cassette comprising the sequence set forth in SEQ ID NQ:5 which codes for said myosin light chain 1/3 enhancer, said CMV promoter and said cDNA sequence of the human serum albumin.

In general, the transcriptional control sequence further comprises at least one sequence responsible for the polyadenylation of the transcribed RNA molecules. Said sequence may be selected in the group consisting of bGH (bovine growth hormone) polyadenylation signal (EP 173552), 5V40 polyadenylation signal, globine polyadenylation signal.

The invention also concerns an expression cassette further comprising a polynucleotide sequence encoding at least one polypeptide of interest which is distinct from human serum albumin, said polypeptide of interest being co-expressed with human serum albumin in said mammalian cell. It further concerns an expression cassette wherein said polynucleotide sequence is localized downstream of said cDNA sequence, is operably linked to the same transcriptional control sequence thereof and said cDNA and polynucleotide sequences being linked by an additional sequence comprising at least one ribosome binding site that facilitates translation of the encoded messenger. According to a preferred embodiment, said polypeptide of interest is a selectable marker such as an antibiotic resistance polypeptide or an hybrid polypeptide thereof. In a preferred embodiment, said antibiotic is bleomycin or the polynucleotide sequence encoding said hybrid is an hybrid gene bleomycin:thymidine kinase. The thymidine kinase gene encodes a product which is able in conjunction with a pro-drug such as gancyclovir (GCV) to kill cells expressing said gene product.

From yet another perspective, the invention includes a recombinant DNA compound comprising at least one of the expression cassette above described. Said recombinant DNA compound can further include at least one nucleotide sequence containing or expressing targeting sequences, transport sequences, sequences involved in replication or integration, or sequence encoding a selectable marker, for example for antibiotic resistance (ampicillin, phleomycin, chloramphenicol), useful for selecting a cell in which said compound has been introduced. Example of such sequences have been reported in the literature and can be readily obtained by those skilled in the art. The recombinant DNA compound can also be modified in order to be stabilized with specific components such as spermine.

As previously stated, the recombinant DNA compound of the invention may be a vector selected from the group consisting of plasmid vector, viral vector or synthetic vector.

The invention further concerns a host cell transformed with a said recombinant DNA compound. Said host cell is preferably a mammalian cell, and more preferably a human muscular cell. This cell can originate from various tissues including those of muscle, skin, nose, lung, liver, spleen, bone marrow, thymus, heart, lymph, bone, cartilage, pancreas, kidney, gall bladder, stomach, intestine, testis, ovary, uterus, rectum, nervous system, eye, gland, connective tissue, blood, tumor etc.

According to the invention, "transformed" means either transfection or infection, and more broadly designate any transferring step resulting in introduction of said recombinant DNA compound in said host cell. Said transferring step can be implemented by any of a wide variety of ways, including method selected from the group consisting of adenoviral infection, transfection with nucleic acid coated particles such as lipoplexes (cationic lipid /nucleic acid complexes) or polyplexes (cationic polymer /nucleic acid complexes) or the like, calcium phosphate transfection of plasmid, transfection with naked nucleic acid, electroporation method or any combination thereof. However, the particular method for introducing the foreign nucleic acid sequence is not crucial to the invention.

Moreover according to a specific embodiment, said transformed host cell is a human muscle and is further encapsulated. Cell encapsulation methodology has been previously described which allows transplantation of encapsulated cells in treatment of Parkinson's disease (Tresco et al., 1992, ASAIO J., 38, 17-23) or Amyotrophic lateral sclerosis (Aebischer et al., 1996, Hum. Gene Ther., 7, 851-860). According to said specific embodiment, transformed cells are encapsulated by compounds which form a microporous membrane, and said encapsulated cells can further be implanted *in vivo*. Capsules, for example approximately 1 cm in length containing the cells of interest may be prepared employing a hollow microporous membrane fabricated from poly-ether-sulfone (PES) (Akzo Nobel Faser AG, Wuppertal, Germany; Deglon et al, 1996, Hum. Gene Ther., 7, 2135-2146). This membrane has a molecular weight cutoff greater than 1,000,000Da, which permits the free passage of proteins and nutrients between the capsule interior and exterior, while preventing the contact of transplanted cells with host cells. The entrapped cells may be implanted by intradermal, subdermal, intravenous, intramuscular, intranasal, intracerebral, intratracheal, intraarterial, intraperitoneal, intravesical, intrapleural, intracoronary or intratumoral ways. In case where said transformed host cell is a myoblast, it can migrate from the site of injection to muscles where expression of human serum albumin can occur.

The present invention concerns the specific use of a cDNA sequence of the human serum

albumin, of an expression cassette, of a recombinant DNA compound or of a transformed host cell as previously disclosed for the preparation of therapeutic composition intended for gene transfer, especially for the treatment of the human or animal body. Administration into vertebrate target tissues, and more specifically into the muscle, can be performed by different delivery routes (systemic delivery and targeted delivery). According to the present invention, the prepared therapeutic composition is preferably administered into muscle, however prepared therapeutic composition administration can also occur in other tissues of the vertebrate body including those of muscle, skin, brain, lung, liver, spleen, bone marrow, thymus, heart, lymph, bone, cartilage, pancreas, kidney, gall bladder, stomach, intestine, testis, ovary, uterus, rectum, nervous system, eye, gland, connective tissue, blood, tumor. The human serum albumin can thus be excreted in body fluids (eg. blood) allowing its delivery in patient organs or said polynucleotide can be associated with targeting molecules which are capable to point its uptake into targeted cells. Gene therapy literature provides many mechanisms for efficient and targeted delivery and expression of genetic information within the cells of a living organism (see for example European patent application n^o98/401108.0). Said administration may be made by intradermal, subdermal, intravenous, intramuscular, intranasal, intracerebral, intratracheal, intraarterial, intraperitoneal, intravesical, intrapleural, intracoronary or intratumoral injection, with a syringe or other devices. Transdermal administration is also contemplated, as are inhalation or aerosol routes. Intramuscular administration route is preferred.

The present invention allows repeated administrations to the patient without major risk of the administered preparation to induce a significant immune reaction. Administration may be by single or repeated dose, once or several times after a certain period of time. Repeated administration allows a reduction in the dose of recombinant DNA compound administered at a single time. The route of administration and the appropriate dose vary in function of several parameters, for example the individual patient, the side effects of the disease, or the albumin level before treatment.

In general, the concentration of polynucleotide in the pharmaceutical compositions is from about 0.1 µg/ml to about 20 mg/ml.

The active dose, or the amount of recombinant DNA compound which should be injected for obtaining satisfactory serum albumin levels, is from about 1 mg to about 50 mg, preferably from about 2 mg to about 24 mg. Said active dose can be administered in a unique administration or in multiple ones distributed into one or more days. For example, the administration cycle is composed of three days treatment : injection of 2 to 8 mg the first day, 2 to 8 mg the second day and 2 to 8 mg the third day. Preferably, the maximum single dose is 8 mg of DNA administered. The separate administrations can be performed by different delivery routes (systemic delivery and targeted delivery, or targeted deliveries for example). In a preferred embodiment, each should be done into the

same target tissue and most preferably by injection.

In a further preferred embodiment the invention pertains to a pharmaceutical preparation comprising at least one of the complexes described above and also incorporating at least one adjuvant capable of improving the transfection capacity of said complex or gene expression into cell and selected from the group consisting of chloroquine, protic compounds such as propylene glycol, polyethylene glycol, glycerol, ethanol, 1-methyl L-2-pyrrolidone or derivatives thereof, aprotic compounds such as dimethylsulfoxide (DM50), diethylsulfoxide, di-n-propylsulfoxide, dimethylsulfone, sulfolane, dimethyl-formamide, dimethylacetamide, tetramethylurea, acetonitrile or derivatives. The composition may also advantageously comprise a source of a cytokine which is incorporated in the form of a polypeptide or as a polynucleotide encoding the cytokine. Preferably, said cytokine is interleukin 10 (IL-b) (see European patent application, filing number 98 40 1667.5 or related foreign application). The therapeutic composition can further comprise an nuclease inhibitor such as actine G (see European patent application, filing number 98 40 1108.0 or related foreign application), or specific magnesium concentration (see European patent application, filing number 98 40 2424.0 or related foreign application)

Therefore, the invention further provides a pharmaceutical composition comprising the above cited recombinant DNA compounds which may also comprise a pharmaceutically acceptable injectable carrier. The carrier is preferably isotonic, hypotonic or weakly hypertonic and has a relatively low ionic strength, such as provided by a sucrose solution. It includes any relevant solvent, aqueous or partly aqueous liquid carrier comprising sterile, pyrogen-free water, dispersion media, coatings, and/or equivalents. The pH of the pharmaceutical preparation is suitably adjusted and buffered.

In the case of *in vivo* treatment according to the invention, in order to improve the transfection rate, the patient may undergo a macrophage depletion treatment prior to administration of the pharmaceutical preparations described above. Such a technique is described in the literature (refer particularly to Van Rooijen et al., 1997, TibTech, 15, 178-184). The patient can also be pre-treated with prednisolone or equivalent.

The invention concerns a method for treating a disorder associated with hypoalbuminaemia in a mammalian organism comprising the step of transferring a recombinant DNA compound into tissue or cells of said organism, wherein said recombinant DNA compound comprises an expression cassette recited in the present invention and such that human serum albumin is expressed resulting in the treatment of said disorder. However, the methods described in this invention can also be applied to the treatment of other conditions and diseases of the human

and animal body in which there is a failure or relative failure of the liver to produce proteins or sufficient quantities of the said proteins to maintain normal function of the tissues or body. Such proteins would include coagulation factors, thrombopoietin. The methods described in the invention would also apply to conditions or diseases of the human or animal body in which there is a failure of the body or organs of the body to produce sufficient hormones, enzymes, growth factors, cytokines or other substances necessary for normal function or homeostasis. The other proteins that could be used in this invention include, but are not restricted to, erythropoietin for the correction of anaemia, thrombopoietin for the correction of thrombocytopenia, protein C and protein F for patients requiring anticoagulation, and leptin for the treatment of obesity. In addition, the invention can be used to produce hormones and growth factors used by the human body at sites other than the physiological source of such hormones and growth factors. These would include but are not restricted to insulin for the treatment of diabetes, endothelial nitric oxide synthetase (ENOS) for the prevention of thrombosis after coronary artery surgery or coronary balloon dilatation or myocardial ischaemia, Granulocyte Colony Stimulating Factor (GCSF), Macrophage Colony Stimulating Factor (MCSF) or Granulocyte Macrophage Colony Stimulating Factor (GMCSF), and/or interferon for the treatment of hepatitis or cancer.

The following examples illustrate the invention.

Example 1

Case Nos 1, 2 and 3 demonstrate the rise of serum albumin in 3 patients following the administration of plasmid encoding albumin. Each patient received 1.5mg intra-muscular DNA daily for 4 consecutive days.

Case 1: A middle-aged male patient with long-standing cirrhosis due to hepatitis. The patient had marked ascites and a serum albumin level of 1.9mmol/l (normal >3.2mmol/l). The DNA/Promotor was administered once daily, at a dose level of 1.5mg, for four consecutive days. The patient was receiving no additional therapy. Serum albumin level rose to 2.7mmol/l within three weeks and this level was sustained for ten weeks. After sixteen weeks the serum albumin level had fallen to 2.3mmol/l. During this period the ascites improved significantly.

Case 2: An elderly male patient with hepatocellular carcinoma resulting from hepatitis and cirrhosis. The patient was not receiving any therapy for the condition. The serum albumin pre-treatment was 3.0mmol/l. Intra-muscular injections of DNA/Promotor were given, as in case 1. Serum albumin levels rose to 3.9mmol/l within three weeks and remained at normal levels for three months.

Case 3: A patient with cirrhosis due to hepatitis C infection. The patient was receiving diuretics (aldactone and lasix) as treatment for ascites. Pre-treatment the patient's serum albumin level was 2.9mmol/l. This level rose to 3.7mmol/l after the course of treatment with the DNA/Promotor. This was accompanied by a significant improvement in the ascites.

Seq ID NQ:1

11

NheI (7)

EcoRI (1)

1 GAATTCGCTAGCTCGACATGATAAGATACATTGATGAGTTTGGACAAACCAAACTAGCAATCGAATGAAAAAATGCTT
80 TATTTCTGAAATTTCTATGCTATGCTTATTTGTAGCAATTATAAGCTGCACTAAATCTTAAGCAAGCAATTTG
159 ATTATGATATGTTTAAAGTTAAATGCAATGCTATGAGCAAGTTTAAAGCAATTAAGCTTAAAGCAATGCTGCTAGAT

SwaI (239)

238 CATTTGATGTTTATTAAGAACATGTGAGCAAAAGGCCAGCAAAAGGCCAGGAACCGTAAAAAGGCCGCTTGCTGGCG
317 TTTTTCATAGGCTCCGCCCCCTGACGAGCATCACAAAAATCGACGCTCAAGTCAGAGGTGSCGAAACCCGACAGGAC
396 TATAAAGATACCAGGCGTTTCCCCCTGGAAGCTCCCTCGTGCCTCTCCTGTTCCGACCTGCGGCTTACCGGATACCT
475 GTCCGCTTTCTCCCTTCGGGAAGCGTGGCGCTTTCTCAATGCTCACGCTGTAGGTATCTCAGTTCGGTGTAGGTCGT
554 CGCTCCAAGCTGGGCTGTGTGCACGAACCCCCGTTACGCCCCGACCGCTGCGCCTTATCCGGTAACTATCGTCTTGAGT
633 CCAACCCGGTAAGACACGACTTATCGGCACTGGCAGCAGCCACTGGTAACAGGATTAGCAGAGCGAGGTATGTAGGCGG
712 TGCTACAGAGTTCTTGAAGTGGTGGCCTAACTACGGCTACACTAGAAGAACAGTATTTGGTATCTGCGCTCTGCTGAAG
791 CCAGTTACCTTCGGAAAAAGAGTTGGTACCTCTTGATCCGGCAAAACAAACCACCGCTGGTAGCGGTGGTTTTTTTGT
870 GCAAGCAGCAGATTACGCGCAGAAAAAAGGATCTCAAGAAGATCCTTTGATCTTTTCTACGGGGTCTGACGCTCAGTG
949 GAACGAAAACCTACGTTAAGGGATTTTGGTCATGGCTAGTTAATTAAGCTGCAATAAACAATCATTATTTTCATTGGAT
1028 CTGTGTGTTGGTTTTTTGTGTGGGCTTGGGGGAGGGGAGGCCAGAATGACTCCAAGAGCTACAGGAAGGCAGGTGAGA
PvuI (117)
SgfI (117)

1107 GACCCCACTGGACAAACAGTGGCTGGACTCTGCACCATAACACACAATCAACAGGGGAGTGAGCTGGATCGAGCTGCGA

AseI (1190)

1186 TCGCGGCGGCCCACTAGTTAACTCATGAGCGGATACATATTTGAATGTATTTAGAAAAATAAACAAATAGGGGTTCGC
1265 GCACATTTCCCCGAAAAGTGCCACCTGACGTCGACGGATCGGGAGATCCGCTATTAATCCAGAGCCCTTGAAGCCAG
1344 AGGAAGATGTATCTCTGAGTTTGAGGCTACCCTACTCTACAGAAAGAGTTCAGGACAGACATTACACGAGAAGCCCTG
1423 CCCCCTCTCTAAAATAAAAGTATTTTCAGAAGCATAAAGGTCACAGTGTAGAGAAAATGACTGCTACACGTAGTCTTAA

BsrGI (1540)

1502 TTATAGAGGGCTCTTTTTTTTTTTTTTTTGTATCTGTGGTGATCATGTCTTTACATTTTTTTCAAGATAGAAAAGCATGA
1581 TGTCTGTGCGGTATAAATTGTTCTGTTTTGAGCCTGTGTATAACGCTTCTCTCAAGATTTTATAATAGTGCTTTAAC
1660 TGTCCCCACGGGCTAACTTCAGCACACTGTCATGGGACCTAACCTTATTAAATTACCATGTGTGAACCGCTCATAACTC

FspI (1766)

PvuII (1807)

1739 AAGTCGACAGAGGTGCAAAAATGGAGCTGCGCAGGCAGAAAGAGTGATCGTCATTTTTTAAATCCCCACAGCTGGCGAA

SspI (1882)

1818 GCAACAGGTGCCTAATTCCTCATCTTTTAAAAATAACTTTTCAAAGCCTGTGCTGTATAAGCAAATATTTTCAAGTTT
1897 GTTTTTAAACCATCTTCAAGTTACCTTCTCACAAAATACATTATGTGCTGATTTTTTTGTCTCAAAATGACATTTGA

976 AGTCTAAGCATATAAAAAATTTATTTCTTTTTAGAAATGAAATTATTATTTAACTGGAGACTTAAATTGTGTCTTAACCT
 2055 TTGCTCCTCCCTTTTCCCTTTTGTCCCTTCTCTCCCACTCCCTCCCTTCTCTTACATGCTCATGGCGGGCTCT
 2134 TCTCTTTCCTACTCTTCTTCTTCTCTCATCCCTCTCCCTGTCTTGTCTTCACTAAACCTTCCACATGGAAAAA
 2213 TAAATTGTATCTTAAAGCTCGGATCTCCCGATCCCTATGGTGCACTCTCAGTACAATCTGCTCTGATGCCGCATAGTT
 2292 AAGCCAGTATCTGCTCCCTGCTTGTGTGTGGAGGTCGCTGAGTAGTGCGCGAGCAAAATTTAAGCTACAACAAGGCAA
 2371 GGCTTGACCGACAATTGCATGAAGAATCTGCTTAGGGTTAGGCGTTTTGCGCTGCTTCG

2450

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3003

3082

CTCTTCTGTCAACCCACGCCTTTGGCACAATGAAGTGG

1 MetLysTrp

3161 GTAACCTTTATTTCCCTTCTTTTTCTCTTTAGCTCGGCTTATCCAGGGGTGTGTTTCGTGAGATGCACACAAGAGTG
 4 Val Thr Phe Ile Ser Leu Leu Phe Leu Phe Ser Ser Ala Tyr Ser Arg Glu Val Phe Arg Asp Ala His Lys Ser G

3240 AGTTGCTCATCGTTTAAAGATTTGGGAGAAGAAAATTTCAAAGCCTTGGTGTGATTGCCTTTGCTCAGTATCTTCA
 30 Lu Val Ala His Arg Phe Lys Asp Leu Glu Glu Asn Phe Lys Ala Leu Val Leu Ile Ala Phe Ala Glu Tyr Leu Glu

PvuII

3319 GCAGTGTCCATTTGAAGATCATGTAAATTAGTGAATGAAGTAACTGAATTTGCAAAAACATGTGTAGCTGATGAGTCA
 56 nGln Cys Pro Phe Glu Asp His Val Lys Leu Val Asn Glu Val Thr Glu Phe Ala Lys Thr Cys Val Ala Asp Glu Ser
 3398 GCTGAAATTTGTGACAAATCACTTCATACCTTTTTGGAGACAAATTATGCACAGTTGCAACTCTTCGTGAAACCTATG
 83 Ala Glu Asn Cys Asp Lys Ser Leu His Thr Leu Phe Glu Asp Lys Leu Cys Thr Val Ala Thr Leu Arg Glu Thr Tyr G
 3477 GTGAAATGGCTGACTGCTGTGCAAAACAAGAACCTGAGAGAAATGAATGCTTCTTGCAACACAAAGATGACAACCCAAA
 109 Iy Glu Met Ala Asp Cys Cys Ala Lys Glu N Glu Pro Glu Arg Asn Glu Cys Phe Leu Glu His Lys Asp Asp Asn Pro As
 3556 CCTCCCCGATTGGTGAGACCAGAGTTGATGTGACTGCTTTTCATGACAATGAAGAGACATTTTGAAGAAAA
 135 nLeu Pro Arg Leu Val Arg Pro Glu Val Asp Val Met Cys Thr Ala Phe His Asp Asn Glu Glu Thr Phe Leu Lys Lys
 3635 TACTTATATGAAATTGCCAGAAGACATCCTTACTTTATGCCCGGAACCTCTTTCTTTGCTAAAAGGTATAAAGCTG
 162 Tyr Leu Tyr Glu Ile Ala Arg Arg His Pro Tyr Phe Tyr Ala Pro Glu Leu Leu Phe Phe Ala Lys Arg Tyr Lys Ala Ala
 3714 CTTTACAGAATGTTGCCAAGCTGCTGATAAAGCTGCCTGCCTGTTGCCAAGCTCGATGAACCTCGGGATGAAGGGAA
 188 Ala Phe Thr Glu Cys Cys Glu Ala Ala Asp Lys Ala Ala Cys Leu Leu Pro Lys Leu Asp Glu Leu Arg Asp Glu Glu Lys
 3793 GGCTTCGTCTGCCAACAGAGACTCAAATGTGCCAGTCTCCAAAATTTGGAGAAAGAGCTTTCAAAGCATGGGCGATG
 214 sAla Ser Ser Ala Lys Glu N Arg Leu Lys Cys Ala Ser Leu Glu N Lys Phe Glu Glu Arg Ala Phe Lys Ala Trp Ala Val
 3872 GCTCGCTGAGCCAGAGATTTCCCAAAGCTGAGTTTGCAGAAGTTCCAAGTTAGTGACAGATCTTACCAAAGTCCACA
 241 Ala Arg Leu Ser Glu N Arg Phe Pro Lys Ala Glu Phe Ala Glu Val Ser Lys Leu Val Thr Asp Leu Thr Lys Val His T
 3951 CGGAATGCTGCCATGGAGATCTGCTTGAATGTGCTGATGACAGGCGGACCTTGCCAAGTATATCTGTGAAAATCAGGA
 267 hr Glu Cys Cys His Glu Asp Leu Leu Glu Cys Ala Asp Asp Arg Ala Asp Leu Ala Lys Tyr Ile Cys Glu Asn Glu As
 4030 TTCGATCTCCAGTAACTGAAGGAATGCTGAAAAACCTCTGTTGGAAAAATCCCACTGCATTGCCGAAGTGGAAAAAT
 293 pSer Ile Ser Ser Lys Leu Lys Glu Cys Cys Glu Lys Pro Leu Leu Glu Lys Ser His Cys Ile Ala Glu Val Glu Asn
 4109 GATGAGATGCCTGCTGACTTGCTTTCATTAGCTGCTGATTTTGTGAAAGTAAGGATGTTTGGAAAACTATGCTGAGG
 320 Asp Glu Met Pro Ala Asp Leu Pro Ser Leu Ala Ala Asp Phe Val Glu Ser Lys Asp Val Cys Lys Asn Tyr Ala Glu Ala
 4188 CAAAGGATGTCTTCTGGCATGTTTTGTATGAATATGCAAGAAGGCATCCTGATTACTCTGTCTGCTGCTGCTGAG
 346 Ala Lys Asp Val Phe Leu Glu Met Phe Leu Tyr Glu Tyr Ala Arg Arg His Pro Asp Tyr Ser Val Val Leu Leu Leu Arg

267 ACTTGCCAAAGACATATGAAACCACTCTAGAGAAGTGCTGTGCCGCTGCAGATCCTCATGAATGCTATGCCAAAGTGTTCT
 372 gLeuAlaLysThr Tyr GluThr Thr LeuGluLysCysCysAlaAlaAlaAspProHisGluCysTyrAlaLysValPhe
 4346 GATGAATTTAAACCTCTTGTGGAAGAGCCTCAGAATTTAATCAAACAAAACCTGTGAGCTTTTAAAGCAGCTTGGAGAGT
 399 AspGluPheLysProLeuValGluGluProGluAsnLeuIleLysGluAsnCysGluLeuPheLysGluLeuGlyGluT
 4425 ACAATTCCAGAATGCGCTATTAGTTCGTTACACCAAGAAAGTACCCCAAGTGTCAACTCCAACCTCTGTAGAGGTCTC
 425 yrLysPheGluAsnAlaLeuLeuValArgTyrThrLysLysValProGluValSerThrProThrLeuValGluValSe
 4504 AAGAAACCTAGGAAAAGTGGGAGCAAAATGTTGTAACATCCTGAAGCAAAAAGAATGCCCTGTGCAGAAGACTATCTA
 451 rArgAsnLeuGlyLysValGlySerLysCysCysLysHisProGluAlaLysArgMetProCysAlaGluAspTyrLeu

Tth111I (4634)

4583 TCCGTGGTCCTGAACCAGTTATGTGTGTTGCATGAGAAAACGCCAGTAAGTGACAGAGTCACAAAATGCTGCACAGAGT
 478 SerValValLeuAsnGluLeuCysValLeuHisGluLysThrProValSerAspArgValThrLysCysCysThrGluS
 4662 CCTTGGTGAACAGGCGACCATGCTTTTTCAGCTCTGGAAGTCGATGAAACATACGTTCCCAAAGAGTTTAAATGCTGAAAC
 504 erLeuValAsnArgArgProCysPheSerAlaLeuGluValAspGluThrTyrValProLysGluPheAsnAlaGluTh
 4741 ATTCACCTTCCATGCAGATATATGCACACTTCTGAGAAGGAGAGACAAATCAAGAAACAACTGCACCTGTTGAGCTT
 530 rPheThrPheHisAlaAspIleCysThrLeuSerGluLysGluArgGluIleLysLysGluThrAlaLeuValGluLeu
 4820 GTGAAACACAAGCCCAAGGCAACAAAAGAGCAACTGAAAGCTGTTATGGATGATTTGCGAGCTTTGTAGAGAAGTGCT
 557 ValLysHisLysProLysAlaThrLysGluGluLeuLysAlaValMetAspAspPheAlaAlaPheValGluLysCysC

Bsu36I (4968)

4899 GCAAGGCTGACGATAAGGAGACCTGCTTTGCCGAGGAGGGTAAAAAACTTGTTGCTGCAAGTCAAGCTGCCTTAGGCTT
 583 ysLysAlaAspAspLysGluThrCysPheAlaGluGluGlyLysLysLeuValAlaAlaSerGluAlaAlaLeuGlyLe
 4978 ATAACATCACATTTAAAGCATCTCAGCCTACCATGAGAATAAGAGAAAGAAATGAAGATCAAAAGCTTATTCATCTG
 609 u

5057 TTTTTCTTTTTCGTTGGTGTAAGCCAACACCTGTCTAAAAAACATAAATTTCTTTAATCATTTTGCTCTTTTCTCT

PvuII (5168)

5136 GTGCTTCAATTAATAAAAAAAAAAAAAAAAAACCAGCTGAGCGCGGTCGCTACCATTACCAGTTGGTCTGGTGTCAATG

BamHI (5216)

5215 GGGATCCTCTAGATTGAGTCGACGTTACTGGCCGAAGCCGCTTGAATAAGGCCGGTGTGCGTTTGTCTATATGTTATT

5294 TTCCACCATATTGCCGTCTTTTGGCAATGTGAGGGCCCGAAACCTGGCCCTGTCTTCTTGACGAGCATTCTAGGGGT

5373 CTTTCCCTCTCGCCAAAGGAATGCAAGGTCTGTTGAATGTCGTGAAGGAAGCAGTTCCTCTGGAAGCTTCTTGAAGAC

BbrPI (5)

5452 AAACAACGTCTGTAGCGACCTTTGCAGGACGCGGAACCCCCCCTGGCGACAGGTGCCTCTGCGGCCAAAAGCCACG

5531 TGTATAAGATACACCTGCAAAGGCGGCACAACCCCACTGCCACGTTGTGAGTTGGATAGTTGTGGAAAGAGTCAAAATGG

5610 CTCTCCTCAAGCGTATTCAACAAGGGGCTGAAGGATGCCCAGAAGGTACCCCATTTGATGGGATCTGATCTGGGGCCTC

5689 GGTGCACATGCTTTACATGTGTTTAGTCGAGGTAAAAAAACGTCTAGGCCCCCGAACCACGGGGACGTGGTTTTCTCT

KasI (5818)

5768 TTGAAAAACACGATAATACCATGGtaagtgatctactagttgtgaccggcgccctagtgttgacaattaatcatcgg

5847 catagtatatcggcatagtataatcgaactcactataggaggggccaccatgtcgactactaaccttcttctcttctccta

BslWI (60)

5926 cagCTTCGTACCCCGGCCATCAACACGCGTCTGCGTTCGACCAGGCTGCGCGTTCTCGCGGCCATAGCAACCGACGTAC

1 SerTyrProGlyHisGluHisAlaSerAlaPheAspGluAlaAlaArgSerArgGlyHisSerAsnArgArgTh

6005 GCGGTTGCGCCCTCGCCGGCAGCAAGAAGCCACGGAAGTCCGCCCGGAGCAGAAAATGCCACGCTACTGCGGGTTTAT

25 rAlaLeuArgProArgArgGluGluGluAlaThrGluValArgProGluGluLysMetProThrLeuLeuArgValTyr

XcmI (6118)

6084 ATAGACGGTCCCCACGGGATGGGAAAAACCAACACGCAACTGCTGGTGGCCCTGGGTTGCGCGACGATATCGTCT

52 IleAspGlyProHisGlyMetGlyLysThrThrThrThrGluLeuLeuValAlaLeuGlySerArgAspAspIleValT

6163 ACGTACCCGAGCCGATGACTTACTGGCGGGTCTGGGGGCTTCCGAGACAATCGCGAACATCTACACCACACAAACCCG

78 yrValProGluProMetThrTyrTrpArgValLeuGlyAlaSerGluThrIleAlaAsnIleTyrThrThrGluHisAr

SphI (6310)

6242 CCTCGACCAGGGTGAGATATCGGCCGGGACGCGGCGGTGGTAATGACAAGCGCCGAGATAACAATGGGCATGCCTTAT

104 gLeuAspGluGlyGluIleSerAlaGlyAspAlaAlaValValMetThrSerAlaGluIleThrMetGlyMetProTyr

521 GCCGTGACCGACGCCGTTCTGGCTCCTCATATCGGGGGGAGGCTGGGAGCTCACATGCCCCGCCCCGGCCCTCACCC
 131▶ AlaVal ThrAspAlaVal LeuAlaProHisIleGlyGlyGluAlaGlySer SerHisAlaProProAlaLeuThrL
 6400 TCATCTTCGACCGCCATCCCATCGCGCCCTCCTGTGCTACCGGCGCGGTACCTTATGGGCAGCATGACCCCCA
 157▶ euIlePheAspArgHisProIleAlaAlaLeuLeuCysTyrProAlaAlaArgTyrLeuMetGlySerMetThrProGly

BspEI (6548)

6479 GGCCGTGCTGGCGTTCTGGCCCTCATCCCGCCGACCTTGCCCGGCACCAACATCGTGCTGGGGCCCTTCCGGAGGAC
 183▶ nAlaVal LeuAlaPheValAlaLeuIleProProThrLeuProGlyThrAsnIleValLeuGlyAlaLeuProGluAsp
 6558 AGACACATCGACCGCCTGGCCAAACGCCAGCGCCCCGGCGAGCGGCTGGACCTGGCTATGCTGGCTGCGATTCCGCCG
 210▶ ArgHisIleAspArgLeuAlaLysArgGlnArgProGlyGluArgLeuAspLeuAlaMetLeuAlaAlaIleArgArgV
 6637 TTTACGGGCTACTTGCCAATACGGTGCGGTATCTGCAGTGCGGCGGGTCTGGCGGGAGGACTGGGGACAGCTTTCGGG
 236▶ aIleTyrGlyLeuLeuAlaAsnThrValArgTyrLeuGlnCysGlyGlySerTrpArgGluAspTrpGlyGlnLeuSerGly
 6716 GACGGCCGTGGCGCCCCAGGGTGCCGAGCCCCAGAGCAACGCGGGCCACGACCCCATATCGGGGACACGTTATTTACC
 262▶ yThrAlaValProProGlnGlyAlaGluProGlnSerAsnAlaGlyProArgProHisIleGlyAspThrLeuPheThr
 6795 CTGTTTCGGGCCCCGAGTTGCTGGCCCCAACGCGGACCTGTATAACGTGTTTGCCTGGGCCTTGACGCTTTGGCCA
 289▶ LeuPheArgAlaProGluLeuLeuAlaProAsnGlyAspLeuTyrAsnValPheAlaTrpAlaLeuAspValLeuAlaL
 6874 AACGCCTCCGTTCCATGCACGTCTTTATCTGGATTACGACCAATCGCCCGCGGCTGCCGGGACGCCCTGCTGCAACT
 315▶ ysArgLeuArgSerMetHisValPheIleLeuAspTyrAspGlnSerProAlaGlyCysArgAspAlaLeuLeuGlnLe

PshAI (6971)

6953 TACCTCCGGGATGGTCCAGACCCACGTACCAACCCCGGCTCCATACCGACGATATGCGACCTGGCGCGCACGTTTGCC
 341▶ uThrSerGlyMetValGlnThrHisValThrThrProGlySerIleProThrIleCysAspLeuAlaArgThrPheAla
 7032 CGTGAGATGATCAGCGGAGCTAATGGCGTCATGGCCAAGTTGACCACTGCCGTTCCGGTGCTCACCGCGCGGACGTCG
 368▶ ArgGluMetIleSerGlyAlaAsnGlyValMetAlaLysLeuThrSerAlaValProValLeuThrAlaArgAspValA

SmaI (7150)

SgrAI (7175)

7111 CCGGAGCGGTGAGTTCTGGACCGACCGGCTCGGGTTCTCCCGGACTTCGTGGAGGACGACTTCGCCGGTGTGGTCCG
 394▶ IaGlyAlaValGluPheTrpThrAspArgLeuGlyPheSerArgAspPheValGluAspAspPheAlaGlyValValAr
 7190 GGACGACGTGACCTGTTTCATCAGCGCGGTCCAGGACCAAGGTGGTGCCGGAACAACACCTGGCCTGGGTGTGGGTGCGC
 420▶ gAspAspValThrLeuPheIleSerAlaValGlnAspGlnValValProAspAsnThrLeuAlaTrpValTrpValArg
 7269 GGCCTGGACGAGCTGTACGCCGAGTGGTCCGAGGTCTGTCCACGAACCTCCGGGACGCCTCCGGGCCGGCCATGACCG
 447▶ GlyLeuAspGluLeuTyrAlaGluTrpSerGluValValSerThrAsnPheArgAspAlaSerGlyProAlaMetThrG
 7348 AGATCGGCAGCAGCCGTGGGGCGGGAGTTCGCCCTGCGCGACCCGGCCGCAACTGCGTGCACTTCGTGGCCGAGGA
 473▶ IuIleGlyGluGlnProTrpGlyArgGluPheAlaLeuArgAspProAlaGlyAsnCysValHisPheValAlaGluGly

RsrII (7457)

SmaI (7476)

7427 GCAGGACTGACCGACCGCCACCAACACCGCCGGTCCGACGGCGGCCACGGGTCCAG
 499▶ uGlnAsp...

EXHIBITS

SEQ ID NQ:1 Sequence of plasmid showing the construct containing the reporter gene plus the required enhancer sequence, promoter sequence and associated sequences to allow the plasmid to function according to this invention.

Figure 1. Diagrammatic construct of plasmid as defined in SEQ ID NQ:1

Figure 2: Mean changes in serum albumin levels in patients treated with either myosin albumin plasmid vector or CMV albumin plasmid.

Figure 3: Sequential changes in serum albumin levels with time in patients treated with albumin plasmid.

Figure 4: Sequential changes in serum albumin levels with time in patient 3 given two injections at day one and after 27 weeks.



CLAIMS

1. An expression cassette for expressing a human serum albumin into a mammalian cell comprising:
 - (i) a cDNA sequence of the human serum albumin,
 - (ii) a transcriptional control sequence operably linked to said cDNA sequence.
2. Expression cassette of claim 1 wherein said cDNA sequence of the human serum albumin has the nucleotide sequence set forth in SEQ ID NO:1
3. Expression cassette of claim 1 wherein said cDNA sequence of the human serum albumin is the nucleic acid sequence of about 2.0kb in length obtained by enzymatic digestion by KpnI and SalI of the plasmid pILMALB5 having the deposit accession number HSRRB HG226
4. Expression cassette of claims 1-3 wherein said transcriptional control sequence comprises a promoter element selected from a viral promoter.
5. Expression cassette of claim 4 wherein said viral promoter is selected from SV40 early and late promoters, the adenovirus major late promoter, the Rous Sarcoma Virus (RSV) promoter, the Cytomegalovirus (CMV) immediate-early promoter and the Majorintermediate-early (MIE) promoter.
7. Expression cassette of claims 4-6 wherein said transcriptional control sequence further comprises at least one enhancer element.
8. Expression cassette of claim 7 wherein said enhancer element is a myosin light chain enhancer.
9. Expression cassette of claim 8 wherein said myosin light chain enhancer is a myosin light chain 1/3 enhancer.
10. Expression cassette of claim 9 wherein said enhancer has the nucleotide sequence set forth in SEQ ID NO:1
11. Expression cassette of claims 7-10 wherein said transcriptional control sequence comprises at the most three enhancer sequences the sequence of which is identical or

selected independently of one another.

12. Expression cassette of claims 7-11 wherein said enhancer sequence is located upstream of said cDNA sequence of the human serum albumin.
13. Expression cassette of claim 1 having the sequence set forth in SEQ ID NO:1 which codes for said myosin light chain 1/3 enhancer, said CMV promoter and said cDNA sequence of the human serum albumin.
14. Expression cassette of claim 1 further comprising a polynucleotide sequence of interest encoding at least one polypeptide of interest distinct from human serum albumin, said polypeptide of interest being co-expressed with human serum albumin in said mammalian cell.
15. Expression cassette of claim 14 wherein said polynucleotide sequence of interest is operably linked to the transcriptional control of said cDNA sequence and comprises at least one ribosome binding site that facilitates translation of messages of any of said polynucleotide sequence of interest.
16. A recombinant DNA compound comprising at least one of the expression cassette of claims 1-15.
17. A compound of claim 16 which comprises at least one nucleotide sequence containing or expressing a selectable marker useful for selecting a cell in which said compound has been introduced.
18. A compound of claim 17 wherein said nucleotide sequence containing or expressing a selectable marker is an antibiotic resistance gene or an hybrid gene thereof.
19. A compound of claim 18 wherein said antibiotic is bleomycin or said hybrid is bleomycin:thymidine kinase
20. A compound of claims 15-18 wherein said recombinant DNA compound is a vector selected from the group consisting in plasmid vector, viral vector or synthetic vector.
21. A host cell transformed with a compound of claims 15-20.

22. A pharmaceutical composition comprising the compound of claims 15-20.
23. Use of a cDNA sequence of the human serum albumin for the preparation of a pharmaceutical composition intended for gene transfer.
24. Use of an expression cassette of claims 1-14 or of a recombinant DNA compound of claims 15-20 for the preparation of a pharmaceutical composition intended for gene transfer.
25. The use of claims 23-25, wherein said gene transfer is gene therapy for the treatment of the human or animal body.
26. The use of claims 23-25, wherein said pharmaceutical composition is designed to be administered in a form which can be injected by the intramuscular route.
27. A method for treating a disorder associated with *hypoalbuminemia* in a mammalian organism comprising the step of transferring a vector into tissue or cells of said organism, wherein said vector comprises an expression cassette of claims 1-14 such that human serum albumin is expressed resulting in the treatment of said disorder.
28. Method of claim 27, wherein said disorder is selected in the group consisting of but not restricted to cirrhosis, liver failure, primary and secondary liver cancers, bacterial or viral infections, trauma.
29. Method of treating a human or animal subject, which subject is suffering from a failure or malfunctioning of the liver to produce protein(s) resulting in conditions such as but not limited to hypoalbuminaemia, anaemia, thrombocytopenia and coagulation disorders, which comprises administering to the skeletal muscle of said subject an effective amount of a construct or vector according to the preceding claims encoding said functional protein such as albumin, erythropoietin, thrombopoietin and coagulation factors II, VII, IX and XI. In so doing the host muscle becomes an extra-hepatic source of production of said protein(s).
30. A construct according to claim 29 which further comprises an enhancer element.
31. A construct according to any one of the preceding claims wherein the functional protein is erythropoietin.

32. A construct according to any one of the preceding claims wherein the functional protein is thrombopoietin.
33. A construct according to any of the preceding claims where the gene encoding for endorphins, insulin, antisense gene for angiotensin, fibroblast growth factor (FGF), vascular endothelial growth factor (VEGF), angiostatin, endostatin, or thrombospondin is included.
34. A construct according to any of the preceding claims where the plasmid is encoded for interferon, or the gene encoding the envelope protein or core protein of viruses such as, but not exclusively, hepatitis B & C or HIV.
35. A construct according to any of the preceding claims where the plasmid is encoding specific tumour associated antigens including, but not exclusively, carcinoembryonic antigen (CEA), CA125, CA19.9 or prostate specific antigen (PSA).
36. A construct according to any of the preceding claims where the plasmid is encoding amino acid sequences or proteins produced or related to the low density lipoprotein (LDL), very low density lipoprotein (VLDL) or high density lipoprotein (HDL) receptors.
37. A construct according to any of the preceding claims where the plasmid is encoding the gene for Endothelial Nitric Oxide Synthetase (ENOS), or Granulocyte Colony Stimulating Factor (GCSF), Macrophage Colony Stimulating Factor (MCSF) or Granulocyte Macrophage Colony Stimulating Factor (GMCSF).
38. A construct according to any of the preceding claims for use in a method of treatment of the human or animal body.



Application No: GB 9930891.8
Claims searched: 1-26 and 30-38

Examiner: L.V.Thomas
Date of search: 20 September 2000

Patents Act 1977 Search Report under Section 17

Databases searched:

UK Patent Office collections, including GB, EP, WO & US patent specifications, in:

UK CI (Ed.R):

Int CI (Ed.7):

Other: Online: EPODOC, WPI, BIOSIS, MEDLINE, CAS-ONLINE, SCISEARCH, EMBASE

Documents considered to be relevant:

Category	Identity of document and relevant passage	Relevant to claims
X	EP 0771874 A2 (STATE OF ISRAEL) see particularly p.3 ll.15-52	1,2
E, X	WO 99/66054 A2 (GENZYME) see particularly pp.1-4, 16, 29-30 and 39	1,14,16,21-25, 31,38
X	WO 93/03164 A1 (RHONE-POULENC ET AL.) see p.4 l.18 - p.5 l.24 and p.8 ll.16-34	1,2
X	WO 92/22635 A1 (UNIV. OF CONNECTICUT ET AL.) see p.2 l.25 - p.4 l.6 and pp.11-13	1,2
X	Transgenic Research 1994, 3(3), pp.141-151 Barash et al. - see the abstract and "Materials and methods" on pp.142-144	1,2

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